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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
10/537,507	06/03/2005	Steffen Thiel	THIEL3	THIEL3 4712	
1444 BROWDY AN	7590 10/03/2007 ID NEIMARK, P.L.L.C.		EXAMINER		
624 NINTH STREET, NW SUITE 300 WASHINGTON, DC 20001-5303			SITTON, JEHA	ANNE SOUAYA	
			ART UNIT	PAPER NUMBER	
			1634	1634	
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			MAIL DATE	DELIVERY MODE	
			10/03/2007	PAPER	

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

A SECTION OF THE PROPERTY OF T						
	Application No.	Applicant(s)				
	10/537,507	THIEL ET AL.				
Office Action Summary	Examiner	Art Unit				
	Jehanne S. Sitton	1634				
The MAILING DATE of this communication app Period for Reply	pears on the cover sheet with the c	orrespondence address				
A SHORTENED STATUTORY PERIOD FOR REPL WHICHEVER IS LONGER, FROM THE MAILING D. - Extensions of time may be available under the provisions of 37 CFR 1.1 after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory period Failure to reply within the set or extended period for reply will, by statute Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION 36(a). In no event, however, may a reply be time will apply and will expire SIX (6) MONTHS from a cause the application to become ABANDONE	N. nely filed the mailing date of this communication. D (35 U.S.C. § 133).				
Status						
1) Responsive to communication(s) filed on 03 J	<u>une 2005</u> .					
· <u> </u>	, 					
3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is						
closed in accordance with the practice under Ex parte Quayle, 1935 C.D. 11, 453 O.G. 213.						
Disposition of Claims						
4) ⊠ Claim(s) 47-87 is/are pending in the applicatio 4a) Of the above claim(s) is/are withdra 5) □ Claim(s) is/are allowed. 6) □ Claim(s) is/are rejected. 7) □ Claim(s) is/are objected to. 8) ⊠ Claim(s) 47-87 are subject to restriction and/or	wn from consideration.	· .				
Application Papers						
9) The specification is objected to by the Examine	er.					
10) ☐ The drawing(s) filed on is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.						
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).						
Replacement drawing sheet(s) including the correct 11) The oath or declaration is objected to by the Ex	· · · · · · · · · · · · · · · · · · ·	•				
Priority under 35 U.S.C. § 119						
12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of: 1. Certified copies of the priority document 2. Certified copies of the priority document 3. Copies of the certified copies of the priority application from the International Burea * See the attached detailed Office action for a list	ts have been received. Is have been received in Applicati rity documents have been receive u (PCT Rule 17.2(a)).	on No ed in this National Stage				
Attachment(s)						
 Notice of References Cited (PTO-892) Notice of Draftsperson's Patent Drawing Review (PTO-948) Information Disclosure Statement(s) (PTO/SB/08) Paper No(s)/Mail Date	4) Interview Summary Paper No(s)/Mail Di 5) Notice of Informal F 6) Other:	ate				

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DETAILED ACTION

Election/Restrictions

1. Restriction is required under 35 U.S.C. 121 and 372.

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in reply to this action, to elect a single invention to which the claims must be restricted.

Group 1, claim(s) 47 and 58-66, in part, drawn to a method of determining a predisposition to an immune system related disease by detecting a polymorphism in SEQ ID NO: 3, using nucleic acid based detection methods.

Group 2, claim(s) 47 and 58, in part, drawn to a method of determining a predisposition to an immune system related disease by detecting a polymorphism in SEQ ID NO: 4.

Group 3, claim(s) 47-57, in part, in part, drawn to a method of determining a predisposition to an immune system related disease by detecting a polymorphism in SEQ ID NO: 1, using protein based methods.

Group 4, claim(s) 47, 51-57, in part, drawn to a method of determining a predisposition to an immune system related disease by detecting a polymorphism in SEQ ID NO: 2 using protein based methods.

Group 5, claim(s) 69, and 78-81, drawn to an isolated oligonucleotide encoding a MASP-2 variant, SEQ ID NO: 1, as well as gene therapy vectors comprising such nucleic acid.

Group 6, claim(s) 70, drawn to an isolated oligonucleotide encoding a Map-19 variant, SEQ ID NO: 2.

Group 7, claim(s) 72-75, in part, drawn to an antibody capable of recognizing MASP-2.

Group 8, claim(s) 72-75, in part, drawn to an antibody capable of recognizing Map-19.

Group 9, claim(s) 76, drawn to a method of treatment for an immune system related disease comprising administering an effective amount of a polypeptide of SEQ ID NO: 1 and/or SEQ ID NO: 2.

Group 10, claim(s) 77, in part, drawn to a polypeptide of SEQ ID NO: 1.

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Group 11, claim(s) 77, in part, drawn to a polypeptide of SEQ ID NO: 2.

Group 12, claim(s) 82,83, 86 and 87 in part, drawn to a method of treatment for inhibition of activity of the lectin component pathway comprising administering an effective amount of SEQ ID NO: 1.

Group 13, claim(s) 82, 83, 86, and 87 in part, drawn to a method of treatment for inhibition of activity of the lectin component pathway comprising administering an effective amount of SEQ ID NO: 2.

Group 14, claim(s) 82, 84, 86, and 87 in part, drawn to a method of treatment for inhibition of activity of the lectin component pathway comprising administering an effective amount of an oligonucleotide of SEQ ID NO: 3.

Group 15, claim(s) 82, and 85-87, in part, drawn to a method of treatment for inhibition of activity of the lectin component pathway comprising administering an effective amount of an antibody which recognizes MASP-2.

Group 16, claim(s) 82, and 85-87, in part, drawn to a method of treatment for inhibition of activity of the lectin component pathway comprising administering an effective amount of an antibody which recognizes Map-19.

2. Claims 67, 68 and 71 link(s) inventions 5 and 6. The restriction requirement between the linked inventions is **subject to** the nonallowance of the linking claim(s). Upon the indication of allowability of the linking claim(s), the restriction requirement as to the linked inventions **shall** be withdrawn and any claim(s) depending from or otherwise requiring all the limitations of the allowable linking claim(s) will be rejoined and fully examined for patentability in accordance with 37 CFR 1.104 **Claims that require all the limitations of an allowable linking claim** will be entered as a matter of right if the amendment is presented prior to final rejection or allowance, whichever is earlier. Amendments submitted after final rejection are governed by 37 CFR 1.116; amendments submitted after allowance are governed by 37 CFR 1.312.

Applicant(s) are advised that if any claim presented in a continuation or divisional application is anticipated by, or includes all the limitations of, the allowable linking claim, such

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claim may be subject to provisional statutory and/or nonstatutory double patenting rejections over the claims of the instant application. Where a restriction requirement is withdrawn, the provisions of 35 U.S.C. 121 are no longer applicable. *In re Ziegler*, 443 F.2d 1211, 1215, 170 USPQ 129, 131-32 (CCPA 1971). See also MPEP § 804.01.

- 3. The inventions listed as Groups 1-16 do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons: The claims lack a special technical feature over the prior art. For example, claim 67 is drawn to a nucleic acid molecule which is only 10 contiguous nucleotides of SEQ 3. However, the prior art of Brennan (US Patent 5,474,796) teaches arrays of nucleic acids of each possible 10 mer nucleic acid sequence. Accordingly, the claims lack a special technical feature over the prior art and lack unity of invention.
- 4. Applicant is advised that the reply to this requirement to be complete must include (i) an election of a species or invention to be examined even though the requirement be traversed (37 CFR 1.143) and (ii) identification of the claims encompassing the elected invention.

The election of an invention or species may be made with or without traverse. To reserve a right to petition, the election must be made with traverse. If the reply does not distinctly and specifically point out supposed errors in the restriction requirement, the election shall be treated as an election without traverse.

Should applicant traverse on the ground that the inventions or species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the inventions or species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C.103(a) of the other invention.

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5. Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR

1.48(b) and by the fee required under 37 CFR 1.17(i).

6. Any inquiry concerning this communication or earlier communications from the examiner should be directed to examiner Jehanne Sitton whose telephone number is (571) 272-0752. The examiner can normally be reached Monday-Thursday from 8:00 AM to 5:00 PM and on alternate Fridays.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Ram Shukla, can be reached on (571) 272-0735. The fax phone number for this Group is (571) 273-8300.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to (571) 272-0547.

Patent applicants with problems or questions regarding electronic images that can be viewed in the Patent Application Information Retrieval system (PAIR) can now contact the USPTO's Patent Electronic Business Center (Patent EBC) for assistance. Representatives are available to answer your questions daily from 6 am to midnight (EST). The toll free number is (866) 217-9197. When calling please have your application serial or patent number, the type of document you are having an image problem with, the number of pages and the specific nature of the problem. The Patent Electronic Business Center will notify applicants of the resolution of the problem within 5-7 business days. Applicants can also check PAIR to confirm that the problem has been corrected. The USPTO's Patent Electronic Business Center is a complete service center supporting all patent business on the Internet. The USPTO's PAIR system provides Internet-based access to patent application status and history information. It also enables applicants to view the scanned images of their own application file folder(s) as well as general patent information available to the public.

For all other customer support, please call the USPTO Call Center (UCC) at 800-786-9199.

/Jehanne Sitton/ Primary Examiner Art Unit 1634 9/28/2007